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(54) **pH Adjusted nonionic surfactant containing alkaline cleaner composition for cleaning microelectronics substrates**

PH-regulierter alkalischer Reiniger mit nicht ionischem Tensid zur Reinigung mikroelektronischer Substrate

Composition de nettoyage ayant un pH déterminé contenant un agent tensioactif nonionique pour nettoyer les substrats microélectroniques

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(56) References cited:
EP-A- 0 578 507 WO-A-93/14884
GB-A- 1 573 208 US-A- 3 886 099
US-A- 4 239 661 US-A- 4 339 340
US-A- 4 612 058 US-A- 4 833 067

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Description

[0001] This invention relates to cleaners for use in the microelectronics industry for cleaning integrated circuit substrates, more particularly for cleaning wafers and vias, without producing undue etching of wafer surfaces.

[0002] The cleaning of integrated circuit (IC) substrates, such as silicon wafers, with metal-free alkaline solutions to remove organic and metal contamination is widely practiced. One commonly used alkaline solution of this type is known as SC-1 or RCA-1 and comprises an aqueous mixture of ammonium hydroxide, hydrogen peroxide, and water (1:1:5 of 30% H_2O_2 , 28% NH_4OH and H_2O). Various cleaning tasks can be accomplished with SC-1, among these, the cleaning of silicon wafers immediately after their fabrication, the cleaning of such wafers immediately prior to gate oxide growth, the removal of oxide etch residues later in the IC processing sequence, and selective etching and resist particulate removal. The purpose of the hydrogen peroxide is to protect the silicon metal from exposure to strong acids or bases by continuously forming a protective oxide layer in order to prevent etching or roughening of the silicon surface.

[0003] However, the presence of hydrogen peroxide in SC-1 formulations imparts an inherent instability to these solutions. Such solutions typically exhibit peroxide half-lives of less than one hour at 70°C. The hydrogen peroxide in the SC-1 solution in the presence of certain metals, particularly copper and iron, becomes unstable and decomposes in rapid exothermic fashion leading to potentially dangerous conditions. The hydrogen peroxide has a low tolerance for metal contamination. Additionally, the decomposed hydrogen peroxide drops the concentration of the hydrogen peroxide leading to the possibility of silicon etching producing wafers that are not acceptable for IC manufacture. Thus, the decomposed hydrogen peroxide needs to be replenished and this changes the solution composition thereby varying the cleaning properties of the solution. In addition, the inherently high pH of the hydrogen peroxide solution presents undesirable safety and environmental concerns.

[0004] Hydrogen peroxide-free IC substrate cleaning solutions comprising simple aqueous solutions of quaternary ammonium hydroxides have been reported in Japanese Patent Publication No. 63-114132 and in U.S. Patent 5,259,888. Oxide etch residue removal with such solutions has also been disclosed in EP Patent Publication 540,261 and Japanese Patent Publication No. 05-259066. However, as disclosed in U.S. Patents 4,113,551 and 4,339,340, such solutions etch (poly)silicon and single crystal silicon wafers resulting in rough surfaces that may be unsuitable for further processing. During the later stages of IC wafer fabrication, the original polished silicon surface has been covered with various oxide and metal layers. However, the backside of the wafer remains exposed to process fluids and must be protected similarly to the initial polished silicon surface. This silicon etching and resulting surface damage may be prevented by including hydrogen peroxide in the alkaline cleaning formulation as disclosed in Japanese Patent Publication No. 3-93229 and U.S. Patents 4,239,661 and 4,339,340. Other agents that reportedly suppress silicon etching include hydrazine as disclosed in Japanese Patent Publication No. 64-14924 and a certain nonionic surfactant, polyoxyethylene nonylphenol, as disclosed in U.S. Patents 4,239,661 and 4,339,340. However, polyoxyethylene nonylphenol has recently been implicated as an environmental estrogen mimic.

[0005] It is therefore desirable that an alkaline cleaner solution be available which does not require the presence of hydrogen peroxide, that is, a cleaner composition in which the addition of hydrogen peroxide or other oxidizing agents is optional. More particularly, it is an object of this invention to provide such an alkaline cleaner solution that does not require the presence of hydrogen peroxide or other oxidizing agent yet produces an effective wafer cleaning action without producing undue wafer roughness sufficient to render the wafers unacceptable for IC manufacturing and processing, particularly for high density IC manufacturing. A still further object of this invention is to provide such an effective cleaning solution that can be used to clean vias in processed wafers of resist and etching residue, i.e. residual ash and organic and inorganic contaminants. A still further object of this invention is to provide such alkaline cleaning compositions for cleaning wafers and producing a roughness of less than about 2.5 nm as the average distance in the Z direction between wafer peak heights and valleys.

[0006] Alkaline cleaner compositions of this invention are provided by alkaline compositions containing aqueous metal ion free bases, a nonionic surfactant and a sufficient amount of a component to control or reduce the pH of the resulting cleaning solutions to a pH within the range of from pH 8 to pH 10 with the proviso that the nonionic surfactant is not selected from the group consisting of monoalkylethers of di-, tri- or tetraethylene glycols and monoalkylethers of di-, tri-, or tetrapropylene glycols containing from 1 to 4 C-atoms in the alkyl radical, urea, acid esters of phosphoric acid, polyglycol ethers, polyvinyl alcohols and polysaccharide derivatives. It has been found that all three components are required to produce an alkaline cleaner composition having effective wafer cleaning action without producing undesirable wafer surface roughness. As the data in the following examples demonstrates, cleaner compositions containing only the base alone, or the base combined with only the nonionic surfactant, or the base combined with only the pH control component, are unable to produce alkaline cleaning compositions producing effective cleaning action as well as not producing undue wafer surface roughness, i.e. a Z-range roughness of less than 2.5 nm. In further embodiments of this invention, the alkaline cleaning compositions may additionally contain metal chelating agents and/or oxidizing agents and the like.

[0007] The alkaline cleaning compositions of this invention generally comprise an alkaline component in an amount

of up to 25% by weight, generally from 0.1 to 10% by weight, a nonionic surfactant in an amount of up to 5% by weight, generally from 0.01 to 2% by weight, and a sufficient amount of a component to reduce or control the pH of the resulting cleaning solution to a pH within the range of from pH 8 to pH 10, generally said amount being within the range of from 0.1 to 10% and preferably from 0.5 to 2% by weight of the total cleaner composition, the remaining balance of the cleaner composition being made up of water, preferably high purity deionized water.

[0008] Any suitable alkaline component may be used in the cleaner compositions of this invention. The alkaline components of these cleaners are preferably quaternary ammonium hydroxides, such as tetraalkyl ammonium hydroxides (including hydroxy- and alkoxy-containing alkyl groups generally of from 1 to 4 carbon atoms in the alkyl or alkoxy group). The most preferable of these alkaline materials are tetramethyl ammonium hydroxide and trimethyl-2-hydroxyethyl ammonium hydroxide (choline). Examples of other usable quaternary ammonium hydroxides include: trimethyl-3-hydroxypropyl ammonium hydroxide, trimethyl-3-hydroxybutyl ammonium hydroxide, trimethyl-4-hydroxybutyl ammonium hydroxide, triethyl-2-hydroxyethyl ammonium hydroxide, tripropyl-2-hydroxyethyl ammonium hydroxide, tributyl-2-hydroxyethyl ammonium hydroxide, dimethylethyl-2-hydroxyethyl ammonium hydroxide, dimethyldi(2-hydroxyethyl) ammonium hydroxide, monomethyltri(2-hydroxyethyl) ammonium hydroxide, tetraethyl ammonium hydroxide, tetrapropyl ammonium hydroxide, tetrabutyl ammonium hydroxide, monomethyltriethyl ammonium hydroxide, monomethyltripropyl ammonium hydroxide, monomethyltributyl ammonium hydroxide, monoethyltrimethyl ammonium hydroxide, monoethyltributyl ammonium hydroxide, dimethyldiethyl ammonium hydroxide, dimethyldibutyl ammonium hydroxide, and the like and mixtures thereof.

[0009] Other alkaline components are also operable including, for example, ammonium hydroxide, organic amines particularly alkanolamines such as 2-aminoethanol, 1-amino-2-propanol, 1-amino-3-propanol, 2-(2-aminoethoxy)ethanol, 2-(2-aminoethylamino)ethanol, 2-(2-aminoethylamino)ethylamine and the like, and other strong organic bases such as guanidine. Mixtures of these additional alkaline components, particularly ammonium hydroxide, with the aforementioned tetraalkyl ammonium hydroxides are also useful.

[0010] The alkaline cleaner compositions of this invention may contain any suitable nonionic surfactant with the proviso that the nonionic surfactant is not selected from the group consisting of monoalkylethers of di-, tri- or tetraethylene glycols and monoalkylethers of di-, tri-, or tetrapropylene glycols containing from 1 to 4 C-atoms in the alkyl radical, urea, acid esters of phosphoric acid, polyglycol ethers, polyvinyl alcohols and polysaccharide derivatives. Among the various suitable nonionic surfactants useful in the cleaner compositions of this invention there may be mentioned, for example, low foaming nonionic surfactants such as alkynol surfactants, fluorinated surfactants such as fluorinated alkyl alkoxylates such as Fluorad FC-171, fluorinated alkylesters such as FC-430 and FC-431 and fluorinated polyoxyethylene alkanols such as Fluorad FC-170C, aliphatic acid esters of polyhydric alcohols, polyoxyethylene diols, siloxane type surfactants and alkylene glycol monoalkyl ethers such as butoxypropanol with the proviso that the nonionic surfactant is not selected from the group consisting of monoalkylethers of di-, tri- or tetraethylene glycols and monoalkylethers of di-, tri-, or tetrapropylene glycols containing from 1 to 4 C-atoms in the alkyl radical, urea, acid esters of phosphoric acid, polyglycol ethers, polyvinyl alcohols and polysaccharide derivatives. Preferred for use as nonionic surfactants in the alkaline cleaning compositions of this invention are alkyne surfactants, especially 3,5-dimethylhexyne-3-ol (Surfynol-61), fluorinated alkyl polyoxyethylene ethanols, especially Fluorad® FC-170C and alkylene glycol monoalkyl ethers, especially butoxypropanol. The nonionic surfactants help partially to suppress silicon etching and also decrease surface tension of the cleaning compositions resulting in improved wetting of the surface to be cleaned and thereby improving the cleaning action of the composition.

[0011] Any compound or mixture of compounds suitable for reducing the pH of the alkaline cleaner solutions of this invention, and which do not unduly adversely inhibit the cleaning action thereof or interfere with the resulting cleaned wafers, may be employed. As examples of such compounds there may be mentioned, for example, acids, bases and their salts acting as buffers, such as inorganic mineral acids and their salts, weak organic acids having a pKa of greater than 2 and their salts, ammonium salts, and buffer systems such as weak acids and their conjugate bases, for example, acetic acid and ammonium-acetate. Preferred for use as such components are acetic acid, potassium biphthalate, mixtures of ammonium chloride and ammonium acetate, especially a 1:1 mixture of these two salts, and mixtures of acetic acid and ammonia and other amines.

[0012] The cleaning solutions of this invention can be used as is or formulated with additional components such as any suitable metal chelating agents to increase the capacity of the formulation to retain metals in solution. Typical examples of chelating agents for this purpose are the following organic acids and their salts: ethylenediaminetetraacetic acid (EDTA), butylenediaminetetraacetic acid, cyclohexane-1,2-diaminetetraacetic acid, diethylenetriaminepentaacetic acid, ethylenediaminetetrapropionic acid, (hydroxyethyl)ethylenediaminetriacetic acid (HEDTA), methyliminodiacetic acid, propylenediaminetetraacetic acid, nitrilotriacetic acid (NTA), citric acid, tartaric acid, gluconic acid, saccharic acid, glyceric acid, oxalic acid, phthalic acid, maleic acid, mandelic acid, malonic acid, lactic acid, salicylic acid, catechol, 8-hydroxyquinoline, N,N,N',N'-ethylenediaminetetra(methylene-phosphonic acid, and the like.

[0013] Also, the addition of oxidizing agents to these formulations is also beneficial if the removal of organic contamination is an important concern. Hydrogen peroxide and the like is commonly used for this purpose. Any suitable

oxidizing agent may be employed, such as, for example, oxidizing anions, such as, for example, nitric acid and its salts and nitrate, persulfate, periodate, perbromate, perchlorate, iodate, bromate, and chlorate salts of ammonium.

[0014] The invention is illustrated by, but not limited to the following examples. In the examples, the percentages are by weight unless specified otherwise. A † sign in the first column Tables indicates a reference example.

EXAMPLE 1

[0015] Aqueous solutions of tetramethyl ammonium hydroxide (TMAH) comprising 0.1, 1.0, and 10% TMAH were prepared. Silicon wafer samples (<1,0,0> crystal face, polished on both sides) were placed in these solutions for 10 minutes at 90°C. After treatment, the "R_z roughness" (defined as the average distance in the z direction between peak heights and valleys) was measured for each concentration. The pH of each solution was also recorded. These highly alkaline solutions severely damaged the silicon surfaces, as shown by the following results.

TMAH Concentration %	R _z (Angstroms)**	pH
† 0.1	13,500	12.4
† 1.0	14,500	13.1
† 10	20,000	14.2

** 1 Angstrom corresponds to 10⁻¹⁰ m

[0016] The formulations were repeated with the addition of a sufficient amount of a 1:1 mixture of ammonium chloride and ammonium acetate to reduce the pH of the solutions to the indicated pH levels.

TMAH Concentration %	R _z (Angstroms)**	pH
† 0.1	200	8.0
† 0.1	500	7.1
† 1.0	200	9.2
† 1.0	500	8.0
† 10	600	9.4

** 1 Angstrom corresponds to 10⁻¹⁰ m

Some reduction in roughness was achieved by pH reduction in this way. However, the silicon surfaces were also too rough to be useful for further IC processing.

[0017] The formulations were repeated incorporating 0.01%, 0.1%, or 1.0% of the surfactant 3,5-dimethylhexyne-3-ol (Surfynol-61), in the aqueous solutions of TMAH but with no addition of components to reduce the pH of the solutions. Surfactant addition alone gave little improvement in surface roughness as shown by the following data.

TMAH Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
† 0.1	0.01	10,100	12.1
† 0.1	0.1	4,000	12.0
† 0.1	1.0	450	12.1
† 1.0	0.01	15,000	12.9
† 1.0	0.1	30,000	13.0
† 1.0	1.0	4,000	13.2
† 10	1.0	7,600	14.2

** 1 Angstrom corresponds to 10⁻¹⁰ m

[0018] Next, cleaner compositions were prepared comprising aqueous solutions of TMAH, Surfynol-61 surfactant and a sufficient amount of a 1:1 mixture of ammonium chloride and ammonium acetate to adjust the pH of the solutions to the indicated pH levels. The observed results were as follows.

TMAH Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
0.1	0.01	<25	9.0
0.1	0.1	30	8.0
0.1	1.0	<25	8.0
1.0	0.01	40	9.3
† 1.0	0.1	<25	6.4
1.0	0.1	<25	9.2*
1.0	1.0	<25	9.4
10	1.0	<25	8.1

* this run utilized potassium biphthalate for pH adjustment.

** 1 Angstrom corresponds to 10^{-10} m

[0019] The silicon wafers resulting from this series of cleaner treatments, which combine the use of an alkaline component and a nonionic surfactant with effective pH control are of sufficiently low roughness to be suitable for typical integrated circuit manufacturing.

EXAMPLE 2

[0020] Alkaline cleaning solutions were evaluated for their ability to produce smooth silicon surfaces at a lower temperature, 70°C, using the same procedure as in Example 1. The observed results for aqueous TMAH solutions alone were as follows.

TMAH Concentration %	R _z (Angstroms)**	pH
† 0.1	5,800	12.0
† 1.0	9,500	13.1
† 10	6,000	14.2

** 1 Angstrom corresponds to 10^{-10} m

These highly alkaline solutions severely damaged the silicon surfaces even at 70°C.

[0021] The formulations were repeated with the addition of a sufficient amount of a 1:1 mixture of ammonium chloride and ammonium acetate to reduce the pH of the solutions to the indicated pH levels. The results obtained were as follows.

TMAH Concentration %	R _z (Angstroms)**	pH
† 1.0	300	9.2
† 1.0	100	8.0
† 10	300	9.4

** 1 Angstrom corresponds to 10^{-10} m

Some reduction in roughness was achieved by pH reduction in this way. However, the silicon surfaces again remained too rough for typical IC processing.

[0022] The formulations were repeated incorporating 0.01%, 0.1%, or 1.0% of 3,5-dimethylhexyne-3-ol. Surfactant addition alone gave little improvement in surface roughness, as shown by the following data.

TMAH Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
† 0.1	0.01	5,000	12.1
† 0.1	0.1	700	12

** 1 Angstrom corresponds to 10^{-10} m

(continued)

TMAH Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
† 0.1	1.0	75	12.1
† 1	0.1	12,500	13
† 1	1.0	1,200	13.2
† 10	1.0	4,000	14.2

** 1 Angstrom corresponds to 10⁻¹⁰ m

[0023] Next, cleaner compositions were prepared comprising aqueous solutions of TMAH, Surfynol-61 surfactant and a sufficient amount of 1:1 mixture of ammonium chloride and ammonium acetate to adjust the pH of the solutions to the indicated pH levels. The observed results were as follows.

TMAH Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
0.1	0.01	<25	9.0
0.1	0.1	<25	8.0
0.1	1.0	<25	8.0
1.0	0.1	30	9.3
1.0	0.1	<25	9.2*
1.0	1.0	<25	9.4
10	1.0	<25	8.1

* this run utilized potassium biphthalate for pH adjustment.

** 1 Angstrom corresponds to 10⁻¹⁰ m

The silicon wafers resulting from this series of cleaner treatments, which combine the use of an alkaline component and a nonionic surfactant with effective pH control are of sufficiently low roughness to be suitable for typical integrated circuit manufacturing.

EXAMPLE 3

[0024] Alkaline cleaning solutions were evaluated for their ability to produce smooth silicon surfaces at an additional lower temperature, 50°C, using the same procedure as in Example 1. The observed results for aqueous TMAH solutions alone were as follows.

TMAH Concentration %	R _z (Angstroms)**	pH
† 1.0	700	13.1
† 10	700	14.2

** 1 Angstrom corresponds to 10⁻¹⁰ m

These highly alkaline solutions damaged the silicon surfaces even at 50°C.

[0025] The 1.0% TMAH concentration in the cleaner composition was repeated with the addition of a 1:1 mixture of ammonium chloride and ammonium acetate to reduce the pH of the solution to the indicated pH level.

TMAH Concentration %	R _z (Angstroms)**	pH
† 1	500	9.2

** 1 Angstrom corresponds to 10⁻¹⁰ m

Some reduction in roughness was achieved by pH reduction in this way. However, the resulting silicon surface was too rough for typical IC processing.

[0026] These formulations were repeated incorporating 0.01%, 0.1%, or 1.0% of the surfactant 3,5-dimethylhexyne-

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3-ol (Surfynol-61) in the aqueous TMAH solution with no addition of components to reduce the pH level of the solutions. Surfactant addition alone gave little improvement in surface roughness, as shown by the following observed data.

TMAH Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
† 1.0	0.1	2,500	13
† 1.0	1.0	150	13.2
† 10	1.0	1,600	14.2

** 1 Angstrom corresponds to 10⁻¹⁰ m

[0027] Next, cleaner compositions were prepared comprising aqueous solutions of TMAH, Surfynol-61 surfactant and a sufficient amount of a 1:1 mixture of ammonium chloride and ammonium acetate to adjust the pH of the solutions to the indicated pH levels. The observed results were as follows.

TMAH Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
1	0.1	<25	9.3
1	0.1	<25	9.2*
1	1	<25	9.4
10	1	<25	8.1

* this run utilized potassium biphthalate for pH adjustment.

** 1 Angstrom corresponds to 10⁻¹⁰ m

The silicon wafers resulting from this series of cleaner treatments, which combine the use of an alkaline component and a nonionic surfactant with effective pH control are of sufficiently low roughness to be suitable for typical integrated circuit manufacturing.

EXAMPLE 4

[0028] Examples 1-3 demonstrate the use of controlled pH alkaline mixtures containing the nonionic surfactant 3,5-dimethylhexyne-3-ol as cleaning solutions without damaging the surface of silicon wafers. This example illustrates the use of other nonionic surfactants.

[0029] Cleaner compositions of aqueous solutions of TMAH and the indicated surfactants, both without and then with the addition of acetic acid to adjust the pH of the solutions to the indicated pH levels, were prepared and tested on silicon wafer surfaces at 90°C for 10 minutes as described in Example 1. The following data was observed.

TMAH Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
† 1.0	Surfynol-104E (.05)	1,600	13.2
1.0	Surfynol-104E (.05)	<25	9.4
† 1.0	Surfynol-104E (0.1)	500	13.3
1.0	Surfynol-104E (0.1)	<25	9.4
† 1.0	Surfynol-400 (0.1)	250	13.3
1.0	Surfynol-400 (0.1)	<25	9.5
† 1.0	Fluorad® FC-170C (0.1)	950	13
† 1.0	Fluorad® FC-170C (0.1)	<25	7.9*
† 1.0	Fluorad® FC-431 (0.1)	350	13.2
1.0	Fluorad® FC-431 (0.1)	<25	9.3
† 1.0	Silwet L-7604 (0.1)	1,450	13.5

* this example utilized a 1:1 mixture of ammonium chloride and ammonium acetate to adjust the pH of the aqueous solution.

** 1 Angstrom corresponds to 10⁻¹⁰ m

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(continued)

TMAH Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
1.0	Silwet L-7604 (0.1)	<25	9.2
† 1.0	Silwet L-7607 (0.1)	1,150	13.5
1.0	Silwet L-7607 (0.1)	<25	9.3
† 0.5	1-butoxy-2-propanol (2.0)	750	12.6
0.5	1-butoxy-2-propanol (2.0)	<25	9.2

** 1 Angstrom corresponds to 10⁻¹⁰ m

[0030] Only when the pH had been adjusted to within the range of pH 8.0 to 10.0 was the surface of the wafers of sufficiently low roughness to be suitable for typical integrated circuit manufacturing.

[0031] Similar cleaner compositions of aqueous solutions of TMAH and the indicated surfactants, both without and then with acetic acid to adjust the pH of the solutions to the indicated pH levels, were prepared and tested on silicon wafer surfaces at 70°C for 10 minutes.

TMAH Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
† 1.0	Surfynol-104E (.05)	250	13.2
1.0	Surfynol-104E (.05)	<25	9.4
† 1.0	Surfynol-104E (0.1)	250	13.3
1.0	Surfynol-104E (0.1)	<25	9.4
† 1.0	Surfynol-440 (0.1)	100	13.3
1.0	Surfynol-440 (0.1)	<25	9.5
† 1.0	Fluorad® FC-170C (0.1)	50	13
† 1.0	Fluorad® FC-170C (0.1)	<25	7.9*
† 1.0	Fluorad® FC-430 (0.1)	250	13.3
1.0	Fluorad® FC-430 (0.1)	<25	9.4
† 1.0	Silwet L-7607 (0.1)	650	13.5
1.0	Silwet L-7607 (0.1)	<25	9.3
† 0.5	1-butoxy-2-propanol (2.0)	50	12.6
0.5	1-butoxy-2-propanol (2.0)	<25	9.2

* this example utilized a 1:1 mixture of ammonium chloride and ammonium acetate to adjust the pH levels of the aqueous solutions.

** 1 Angstrom corresponds to 10⁻¹⁰ m

[0032] Only when the pH had been adjusted to within the range of pH 8.0 to 10.0 was the surface of the wafers of sufficiently low roughness to be suitable for typical integrated circuit manufacturing.

[0033] The chemical designations of the surfactants employed in this Example 4 are as follows.

Trade Name	Chemical Description
Surfynol-104E	2,4,7,9-tetramethyl-5-decyne-4,7-diol
Surfynol-440	ethoxylated tetramethyldecyldiol
Fluorad® FC-170C	fluorinated alkyl polyoxyethylene ethanol
Fluorad® FC-430	fluorinated alkyl esters
Fluorad® FC-431	fluorinated alkyl esters
Silwet L-7604	organomodified polymethylsiloxane

(continued)

Trade Name	Chemical Description
Silwet L-7607	polyalkyleneoxide modified polydimethylsiloxane

EXAMPLE 5

[0034] Other metal-free aqueous alkaline base components may be utilized in the cleaner compositions of this invention in place of the aqueous TMAH used in the preceding examples. The following data was obtained similarly to Example 1 at 90°C for 10 minutes with Surfynol-61 utilized as the surfactant and acetic acid utilized to adjust the pH of the aqueous solutions to within a pH range of from pH 8.0 to 10.0.

Base Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
† tetraethylammonium hydroxide (1.0)	none	22,000	13.1
† tetraethylammonium hydroxide (1.0)	0.1	12,500	13.2
tetraethylammonium hydroxide (1.0)	0.1	<25	9.2
† choline (1.0)	none	6,200	12.9
† choline (1.0)	0.1	1,100	13.1
choline (1.0)	0.1	<25	9.2

** 1 Angstrom corresponds to 10⁻¹⁰ m

[0035] The following data for cleaner compositions containing other alkaline base components was obtained similarly at 70°C for 10 minutes.

Base Concentration %	Surfactant Concentration %	R _z (Angstroms)**	pH
† tetraethylammonium hydroxide (1.0)	none	5,000	13.1
† tetraethylammonium hydroxide (1.0)	0.1	3,300	13.2
tetraethylammonium hydroxide (1.0)	0.1	<25	9.2
† choline (1.0)	none	4,300	12.9
† choline (1.0)	0.1	250	13.1
choline (1.0)	0.1	<25	9.2
† 2-aminoethanol (2.0)	none	16,000	11.9
† 2-aminoethanol (2.0)	0.1	1,200	12.0
2-aminoethanol (2.0)	0.1	<25	9.2
† guanidine carbonate (2.0)	none	6,400	11.5
† guanidine carbonate (2.0)	0.1	500	11.5
guanidine carbonate (2.0)	0.1	50	9.2

** 1 Angstrom corresponds to 10⁻¹⁰ m

[0036] The data also shows that the benefits of this invention are obtained with aqueous alkaline cleaner compositions containing a variety of alkaline base components.

EXAMPLE 6

[0037] A cleaner bath containing a preferred formulation of this invention was prepared by mixing: 7.6 l of water, 79 gl of tetramethylammonium hydroxide, 7.9 gl of EDTA, 150 gl of 1-butoxy-2-propanol, and 150 gl of ammonium acetate. This cleaner bath was used to clean silicon wafers (a total of 12,560 cm² in area) at room temperature with the unpolished wafer backs exposed. The silicon wafers had been processed in the manner described hereinafter.

[0038] Silicon wafers were obtained that had been processed in the following manner:

- a. metallization with copper/aluminum alloy followed by titanium and then titanium nitride;
- b. coating with a silicon oxide dielectric using chemical vapor deposition;
- c. lithographic patterning of 0.5 to 1.0 microns vias using a photoresist material;
- d. pattern transfer to the dielectric layer using reactive ion etching;
- e. plasma ashing to remove most of the residual photoresist.

[0039] The resulting patterned wafers were contaminated with a considerable amount of residual ash that comprised a mixture of organic materials from the photoresist and inorganic materials from the underlying metal layer that had been partially exposed during step (d). The use of the hereinbefore described cleaner bath formulation as well as the following controlled pH alkaline cleaning solutions, all of which contained water, tetramethyl ammonium hydroxide, a nonionic surfactant (either 3,5-dimethylhexyne-3-ol or 1-butoxy-2-propanol), and a chelating agent (EDTA) and one of the following listed pH reducing components, removed this residual ash without damaging any silicon or metal features that were present.

pH Reducing Component	pH
Ammonium acetate	9.3-10.0
Ammonium acetate, ammonium hydroxide	9-10
Acetic acid, ammonium hydroxide	9.5-10
Acetic acid, 1-amino-2-propanol	9.5-10
Acetic acid, hydrogen peroxide	9.3
Ammonium acetate, hydrogen peroxide	9.5
Ammonium acetate, nitric acid	9.5
Ammonium nitrate	8.9-10.0
Ammonium chloride, ammonium acetate, hydrogen peroxide	9.6-10.0
Ammonium chloride, ammonium acetate, ammonium periodate	9-10
Ammonium chloride, ammonium acetate, ammonium nitrate	9.4-10.0
Ammonium chloride, ammonium acetate, ammonium persulfate	9-10

[0040] Photoresist ash residues were successfully removed. The bath was analyzed for silicon content after use giving <0.2 ppm of dissolved Si demonstrating that the desired cleaning was achieved without etching exposed silicon or silicon dioxide circuit elements.

EXAMPLE 7

[0041] This example demonstrates the metal removal capabilities of a cleaner formulation for the removal of aluminum, copper, and iron from metal-contaminated silicon wafers. Cleaner Formulation A was prepared by dissolving ethylenediaminetetraacetic acid (EDTA), ammonium acetate, tetramethyl ammonium hydroxide, and 3,5-dimethylhexyne-3-ol in deionized water. Cleaner Formulation B was similarly prepared substituting nitrilotriacetic acid (NTA) for the EDTA. Both formulations exhibited a pH of approximately 10. Metal-contaminated wafers were cleaned in these solutions for 10 minutes at 70°C. The wafers were removed from the cleaner, rinsed in deionized water, and dried. The remaining amount of wafer metal contamination was measured by washing the wafers with dilute hydrochloric acid which was then analyzed for aluminum, copper, and iron. The observed results were as follows.

Cleaning Formulation	Aluminum (micrograms/wafer)	Copper (micrograms/wafer)	Iron (micrograms/wafers)
† none	1	1	0.9
Formulation A	0.1	<0.01	0.2
Formulation B	0.06	<0.01	0.1

EXAMPLE 8

[0042] In another embodiment of the present invention an aqueous alkaline cleaner (Formulation C) containing tetramethyl ammonium hydroxide (0.5%), EDTA (0.1%), ammonium chloride (0.3%), ammonium acetate (0.3%), hydrogen peroxide (1.0%), and 3,5-dimethylhexyne-3-ol (0.1%) was directly compared to a conventional SC-1 cleaner containing, by volume, one part concentrated ammonium hydroxide, one part 30% hydrogen peroxide, and five parts of deionized water. Both cleaning solutions were purposely contaminated with 5 micrograms/liter each of aluminum, iron, and nickel, and 10 micrograms/liter of copper introduced as nitrate salts. Silicon wafers were cleaned in these solutions for 10 minutes at 70°C after which they were rinsed in deionized water, and dried. Residual metal contamination on the wafers was then measured using hydrogen fluoride vapor phase decomposition of the native oxide layer of the silicon wafer followed by scanning the wafer surface with a small volume of water. This water was removed and analyzed by inductively coupled plasma analysis with mass spectral detection giving the following results.

Cleaning Formulation	x 10 ¹⁰ atoms/cm ²			
	Aluminum	Copper	Nickel	Iron
† none (untreated wafer)	42	<6	21	72
† SC-1	2,800	<15	5	743
Formulation C	52	<6	<2	35

[0043] These data clearly show the superior cleaning ability of Formulation C versus that of a conventional cleaner for metal removal from silicon wafer surfaces.

EXAMPLE 9

[0044] Silicon wafers were cleaned as in Example 8 and an Atomic Force Microscope (AFM) was used to examine the surfaces for roughness before and after cleaning. Roughness is reported in this example as "mean roughness" (R_a) which is defined as the mean value of the surface relative to the center plane and is calculated using:

$$R_a = \frac{1}{L_y L_x} \int_0^{L_y} \int_0^{L_x} |f(x, y)| dx dy$$

where $f(x, y)$ is the surface relative to the center plane and L_x and L_y are the dimensions of the surface in two dimensions.

[0045] The untreated, polished wafer with its normal covering of native oxide has an R_a of 0.140 nanometers. When this surface is exposed to SC-1 the R_a value increases to 0.185 nanometers. However, exposure to Formulation C beneficially lowers this R_a value to 0.137 nanometers.

Sample	R_a (nanometers)
† none (untreated wafer)	0.140
† SC-1	0.185
Formulation C	0.137

EXAMPLE 10

[0046] Flamed 57 mm. quartz wafers were used which were stored in sealed quartz petri dishes to avoid organic contamination. These wafers were cleaned as in Example 8 and analyzed for organic contamination using plasma chromatography coupled with mass spectroscopy (PC/MS). This technique involves heating to volatilize any adhering organic materials. The volatilized molecules are ionized and separated into identifiable fractions by passing them through a potential gradient. The high sensitivity of PC/MS allows detection of one part of organic material in 10¹³ parts of matrix.

[0047] The untreated wafer was simply rinsed in deionized water for ten minutes at room temperature. The PC/MS spectrum for this untreated wafer had two ion mass peaks (293 and 337 mass units) which are due to the environmen-

tally ubiquitous phthalate esters, common plasticizers used in laboratory equipment. A wafer cleaned as in Example 8 using SC-1, gave a PC/MS spectrum having six new ion mass peaks (300, 335, 371, 411, 436, 533 mass units) indicative of more organic contamination than the untreated control. A wafer cleaned as in Example 8 using a formulation containing TMAH (1%), EDTA (0.1%), ammonium chloride (0.3%), ammonium acetate (0.3%), hydrogen peroxide (7%) and 3,5-dimethylhexyne-3-ol (0.2%), gave a PC/MS spectrum having three ion mass peaks (300, 337 and 372 mass units). This spectrum shows less organic contamination than that indicated for SC-1. Thus, this cleaner formulation of this invention reduces residual volatile organics on this wafer to less than the standard SC-1 treatment. This Example shows that negligible residue was left by the organic components of this formulation insuring that further IC processing can proceed without interference.

Claims

1. An alkaline cleaning solution for microelectronics substrates comprising an aqueous metal ion free base, a nonionic surfactant and an effective amount of a pH reducing chemical component to reduce or control the pH of the cleaning solution to a pH within the range of from pH 8 to pH 10 with the proviso that the nonionic surfactant is not selected from the group consisting of monoalkylethers of di-, tri- or tetraethylene glycols and monoalkylethers of di-, tri-, or tetrapropylene glycols containing from 1 to 4 C-atoms in the alkyl radical, urea, acid esters of phosphoric acid, polyglycol ethers, polyvinyl alcohols and polysaccharide derivatives.
2. An alkaline cleaning solution for microelectronics substrates according to Claim 1 comprising from 0.1% to 25% by weight of an aqueous metal ion free base selected from ammonium hydroxide, alkanolamines, guanidine, quaternary ammonium hydroxides and mixtures thereof, from 0.01% to 5% by weight of a nonionic surfactant selected from the group consisting of alkynol surfactants, fluorinated alkyl alkoxyates, fluorinated alkyl esters, fluorinated polyoxyethylene alkanols, aliphatic acid esters of polyhydric alcohols, polyoxyethylene diols, siloxane surfactants and alkylene glycol monoalkyl ethers and mixtures thereof and from 0.1% to 10% by weight of a pH reducing chemical component to reduce or control the pH of the cleaning solution to a pH within the range of from pH 8 to pH 10 and wherein the chemical component to reduce or control the pH of the cleaning solution is selected from the group consisting of acids, bases and their salts and buffer systems of weak organic acids and conjugate bases.
3. An alkaline cleaning solution according to any one of Claims 1 or 2 wherein the metal ion free base is selected from a tetraalkyl ammonium hydroxide wherein the alkyl group is an unsubstituted alkyl group or an alkyl group substituent with a hydroxy or alkoxy radical.
4. A cleaning solution according to any one of Claims 1 or 2 wherein the metal ion free base is selected from tetramethyl ammonium hydroxide, tetraethyl ammonium hydroxide and trimethyl-2-hydroxyethyl ammonium hydroxide.
5. A cleaning solution according to any one of Claims 1 or 2 wherein the metal ion free base is an alkanolamine or a guanidine compound.
6. A cleaning solution according to any one of claims 1 to 5 wherein the nonionic surfactant is selected from the group consisting of alkynol surfactants, fluorinated polyoxyethylene alkanol surfactants, siloxane surfactants and alkylene glycol monoalkyl ether surfactants.
7. A cleaning solution according to Claim 6 wherein the nonionic surfactant is selected from the group consisting of 3,5-dimethylhexyne-3-ol, a fluorinated polyoxyethylene ethanol and butoxypropanol.
8. A cleaning solution according to any one of Claims 1 to 7 additionally comprising a metal chelating agent.
9. A cleaning solution according to Claim 8 wherein the metal chelating agent is ethylenediaminetetraacetic acid.
10. A cleaning solution according to any one of Claims 1 to 9 additionally comprising an oxidizing agent.
11. A cleaning solution according to Claim 10 wherein the oxidizing agent is selected from hydrogen peroxide, nitric acid and its salts, and the persulfate, periodate, perbromate, perchlorate, iodate, bromate or chlorate salts of ammonium.
12. A cleaning solution according to any one of Claims 1 to 11 wherein the chemical- component to reduce the pH of

the cleaning solution is selected from acetic acid, potassium biphthalate, a mixture of ammonium acetate with ammonium chloride, and a mixture of acetic acid with ammonia.

13. A cleaning solution according to any one of Claims 1, 2 or 9 comprising water, tetramethyl ammonium, hydroxide, 3,5-dimethylhexyne-3-ol, ammonium chloride and ammonium acetate.
14. A cleaning solution according to any one of Claims 1, 2 or 9 comprising water, tetramethyl ammonium hydroxide, acetic acid, ammonia or an alkanolamine and 1-butoxy-2-propanol.
15. Use of a cleaning composition according to any one of Claims 1 to 14 for cleaning a microelectronics wafer substrate and maintaining wafer surface smoothness.
16. Use of a cleaning solution according to any one of Claims 1 to 14 for cleaning vias in a microelectronic wafer substrate.

Patentansprüche

1. Alkalische Reinigungslösung für mikroelektronische Substrate, umfassend eine wässrige metallionenfreie Base, ein nichtionisches Tensid und eine wirksame Menge eines pH-reduzierenden chemischen Bestandteils zum Reduzieren oder Regeln des pH-Wertes der Reinigungslösung auf einen pH-Wert im Bereich von pH 8 bis pH 10, mit der Maßgabe, dass das nichtionische Tensid nicht ausgewählt ist aus Monoalkylethern von Di-, Tri- oder Tetraethylenglykolen und Monoalkylethern von Di-, Tri- oder Tetrapropylenglykolen mit 1 bis 4 C-Atomen im Alkylrest, Harnstoff, Säureestern der Phosphorsäure, Polyglykolethern, Polyvinylalkoholen und Polysaccharid-Derivaten.
2. Alkalische Reinigungslösung für mikroelektronische Substrate nach Anspruch 1, umfassend 0,1 Gew.-% bis 25 Gew.-% einer wässrigen metallionenfreien Base, ausgewählt aus Ammoniumhydroxid, Alkanolaminen, Guanidin, quartären Ammoniumhydroxiden und Gemischen davon, 0,01 Gew.-% bis 5 Gew.-% eines nichtionischen Tensids, ausgewählt aus Alkinol-Tensiden, fluorierten Alkylalkoxylaten, fluorierten Alkylestern, fluorierten Polyoxyethylenalkanolen, aliphatischen Säureestern von Polyalkoholen, Polyoxyethylendiolen, Siloxan-Tensiden und Alkylenglycolmonoalkylethern und Gemischen davon, sowie 0,1 Gew.-% bis 10 Gew.-% eines pH-reduzierenden chemischen Bestandteils zum Reduzieren oder Regeln des pH-Wertes der Reinigungslösung auf einen pH-Wert im Bereich von pH 8 bis pH 10, wobei der chemische Bestandteil zum Reduzieren oder Regeln des pH-Wertes der Reinigungslösung ausgewählt ist aus Säuren, Basen und ihren Salzen sowie Puffersystemen aus schwachen organischen Säuren und konjugierten Basen.
3. Alkalische Reinigungslösung nach einem der Ansprüche 1 oder 2, wobei die metallionenfreie Base ausgewählt ist aus Tetraalkylammoniumhydroxid, in dem der Alkylrest ein unsubstituierter Alkylrest oder ein Alkylsubstituent mit einem Hydroxy- oder Alkoxyrest ist.
4. Reinigungslösung nach einem der Ansprüche 1 oder 2, wobei die metallionenfreie Base ausgewählt ist aus Tetramethylammoniumhydroxid, Tetraethylammoniumhydroxid und Trimethyl-2-hydroxyethylammoniumhydroxid.
5. Reinigungslösung nach einem der Ansprüche 1 oder 2, wobei die metallionenfreie Base ein Alkanolamin oder eine Guanidinverbindung ist.
6. Reinigungslösung nach einem der Ansprüche 1 bis 5, wobei das nichtionische Tensid ausgewählt ist aus Alkinol-Tensiden, fluorierten Polyoxyethylenalkanol-Tensiden, Siloxan-Tensiden und Alkylenglycolmonoalkylether-Tensiden.
7. Reinigungslösung nach Anspruch 6, wobei das nichtionische Tensid ausgewählt ist aus 3,5-Dimethylhexin-3-ol, einem fluorierten Polyoxyethylenethanol und Butoxypropanol.
8. Reinigungslösung nach einem der Ansprüche 1 bis 7, welche zusätzlich ein Metallkomplexierungsmittel umfasst.
9. Reinigungslösung nach Anspruch 8, wobei das Metallkomplexierungsmittel Ethylendiamintetraessigsäure ist.
10. Reinigungslösung nach einem der Ansprüche 1 bis 9, welche zusätzlich ein Oxidationsmittel umfasst.

11. Reinigungslösung nach Anspruch 10, wobei das Oxidationsmittel ausgewählt ist aus Wasserstoffperoxid, Salpetersäure und ihren Salzen, sowie den Persulfat-, Perjodat-, Perbromat-, Perchlorat-, Jodat-, Bromat- oder Chlorsalzen von Ammonium.
- 5 12. Reinigungslösung nach einem der Ansprüche 1 bis 11, wobei der chemische Bestandteil zum Reduzieren des pH-Wertes der Reinigungslösung ausgewählt ist aus Essigsäure, Kaliumbiphthalat, einem Gemisch von Ammoniumacetat mit Ammoniumchlorid und einem Gemisch von Essigsäure mit Ammoniak.
- 10 13. Reinigungslösung nach einem der Ansprüche 1, 2 oder 9, umfassend Wasser, Tetramethylammoniumhydroxid, 3,5-Dimethylhexin-3-ol, Ammoniumchlorid und Ammoniumacetat.
14. Reinigungslösung nach einem der Ansprüche 1, 2 oder 9, umfassend Wasser, Tetramethylammoniumhydroxid, Essigsäure, Ammoniak oder ein Alkanolamin sowie 1-Butoxy-2-propanol.
- 15 15. Verwendung einer Reinigungszusammensetzung nach einem der Ansprüche 1 bis 14 zur Reinigung eines mikroelektronischen Wafersubstrates und Erhaltung der Glätte der Waferoberfläche.
16. Verwendung einer Reinigungslösung nach einem der Ansprüche 1 bis 14 zur Reinigung von Verbindungsbahnen (vias) eines mikroelektronischen Wafersubstrats.
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Revendications

- 25 1. Solution alcaline de nettoyage pour des substrats microélectroniques comprenant une base aqueuse sans ions métalliques, un tensioactif non ionique et une quantité efficace d'un composant chimique réduisant le pH pour réduire ou contrôler le pH de la solution de nettoyage à un pH situé dans le domaine de pH 8 à pH 10 à condition que le tensioactif non ionique ne soit pas choisi dans le groupe consistant en les monoalkyléthers de di-, tri- ou tétraéthylèneglycols et les monoalkyléthers de di-, tri- ou tétrapropylèneglycols contenant de 1 à 4 atomes de carbone dans le radical alkyle, l'urée, les esters acides de l'acide phosphorique, les polyglycoléthers, les poly (alcool vinyliques) et les dérivés de polysaccharides.
- 30 2. Solution alcaline de nettoyage pour des substrats microélectroniques selon la revendication 1, comprenant de 0,1 % à 25 % en masse d'une base aqueuse sans ions métalliques choisie parmi l'hydroxyde d'ammonium, les alcanolamines, la guanidine, les hydroxydes d'ammonium quaternaire et leurs mélanges, de 0,01 % à 5 % en masse d'un tensioactif non ionique choisi dans le groupe consistant en les tensioactifs alcynols, les alcoylates d'alkyle fluorés, les alkylesters fluorés, les polyoxyéthylènealcanols fluorés, les esters d'acides aliphatiques et d'alcools polyfonctionnels, les polyoxyéthylènealcanols fluorés, les tensioactifs siloxane et les monoalkyléthers d'alkylèneglycols et leurs mélanges et de 0,1 % à 10 % en masse d'un composant chimique réduisant le pH pour réduire ou contrôler le pH de la solution de nettoyage à un pH situé dans le domaine de pH 8 à pH 10 et où le composant chimique pour réduire ou contrôler le pH de la solution de nettoyage est choisi dans le groupe consistant en les acides, les bases et leurs sels et les systèmes tampons d'acides organiques faibles et de bases conjuguées.
- 35 3. Solution alcaline de nettoyage selon l'une quelconque des revendications 1 et 2, où la base sans ions métalliques est choisie parmi un hydroxyde de tétraalkylammonium où le groupe alkyle est un groupe alkyle non substitué ou un groupe alkyle substitué par un radical hydroxyle ou alcoxy.
- 40 4. Solution de nettoyage selon l'une quelconque des revendications 1 et 2, où la base sans ions métalliques est choisie parmi l'hydroxyde de tétraméthylammonium, l'hydroxyde de tétraéthylammonium et l'hydroxyde de triméthyl-2-hydroxyéthylammonium.
- 50 5. Solution de nettoyage selon l'une quelconque des revendications 1 et 2, où la base sans ions métalliques est une alcanolamine ou un composé de la guanidine.
- 55 6. Solution de nettoyage selon l'une quelconque des revendications 1 à 5, où le tensioactif non ionique est choisi dans le groupe consistant en les tensioactifs alcynols, les tensioactifs polyoxyéthylènealcanols fluorés, les tensioactifs siloxane et les tensioactifs monoalkyléthers d'alkylèneglycols.
7. Solution de nettoyage selon la revendication 6, où le tensioactif non ionique est choisi dans le groupe consistant

en le 3,5-diméthylhexyn-3-ol, un polyoxyéthylèneéthanol fluoré et le butoxypropanol.

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8. Solution de nettoyage selon l'une quelconque des revendications 1 à 7, comprenant en outre un agent chélatant les métaux.
9. Solution de nettoyage selon la revendication 8, où l'agent chélatant les métaux est l'acide éthylènediaminetétraacétique.
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10. Solution de nettoyage selon l'une quelconque des revendications 1 à 9, comprenant en outre un agent oxydant.
11. Solution de nettoyage selon la revendication 10, où l'agent oxydant est choisi parmi le peroxyde d'hydrogène, l'acide nitrique et ses sels, et les sels persulfate, periodate, perbromate, perchlorate, iodate, bromate ou chlorate d'ammonium.
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12. Solution de nettoyage selon l'une quelconque des revendications 1 à 11, où le composant chimique pour réduire le pH de la solution de nettoyage est choisi parmi l'acide acétique, le biphthalate de potassium, un mélange d'acétate d'ammonium et de chlorure d'ammonium et un mélange d'acide acétique et d'ammoniac.
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13. Solution de nettoyage selon l'une quelconque des revendications 1, 2 et 9, comprenant de l'eau, de l'hydroxyde de tétraméthylammonium, du 3,5-diméthylhexyn-3-ol, du chlorure d'ammonium et de l'acétate d'ammonium.
14. Solution de nettoyage selon l'une quelconque des revendications 1, 2 et 9, comprenant de l'eau, de l'hydroxyde de tétraméthylammonium, de l'acide acétique, de l'ammoniac ou une alcanolamine et du 1-butoxy-2-propanol.
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15. Utilisation d'une composition de nettoyage selon l'une quelconque des revendications 1 à 14, pour nettoyer un substrat de type plaquette microélectronique et pour maintenir le lissé de la surface de la plaquette.
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16. Utilisation d'une solution de nettoyage selon l'une quelconque des revendications 1 à 14, pour nettoyer des traversées dans un substrat de type plaquette micro-électronique.

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